

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: June 24, 1999, 01:22:16 ; Search time 67.43 Seconds
(without alignments)
410.133 Million cell updates/sec

Title: US-09-205-015-2

Perfect score: 147

Sequence: 1 agataactggccaacatg.....ctcccctctgtttatct 147

Scoring table: IDENTITY_NUC

Searched: 240622 seqs, 94065609 residues

Database : N_Geneseq_34:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Match	Score	Length	ID	Description
1	27	18.4	1420	1 V32475	Bovine retinaldehy
2	26.8	18.2	2158	1 Q25156	Alpha-GalNac from
3	26.8	18.2	1840	1 Q81826	Alpha-N-acetylgluc
c 4	26.8	18.2	334	1 T26078	Human gene signatu
5	26.4	18.0	495	1 N91109	Human reg cDNA., N
6	26.4	18.0	441	1 Q05622	Sequence encoding
7	26.4	18.0	498	1 N81962	Sequence of human
c 8	26.4	18.0	6327	1 T32301	Dermatomyositis sp
9	26.2	17.8	1827	1 T09865	First intron promo
c 10	26	17.7	142	1 V41229	Prevotella nigresc
c 11	25.8	17.6	2525	1 V02016	DNA encoding tumou
c 12	25.6	17.4	383	1 O61216	Human brain Expre
c 13	25.6	17.4	1254	1 O58718	Human gamma-1 chai
c 14	25.4	17.3	1371	1 O47076	B. canis 21B4/rhop
c 15	25.2	17.1	565	1 M70930	Sequence encoding
c 16	25.2	17.1	495	1 Q40477	MURF-1 coding reg
c 17	25.2	17.1	3969	1 Q49757	PTK gene HprtK5. Ne
c 18	25.2	17.1	3969	1 T03099	Protein tyrosine-k
c 19	25.2	17.1	4290	1 Q92641	Human non-differen
c 20	25.2	17.1	4290	1 T18394	Receptor type tyro
c 21	25.2	17.1	4290	1 T42593	Coding sequence fo
c 22	25.2	17.1	4290	1 T51235	Receptor-type tyro
c 23	25.2	17.1	1228	1 T94471	Human Fchd605 gene
c 24	25	17.0	3252	1 Q54375	Human BDNF vector.
c 25	25	17.0	3370	1 Q79534	Bovine tracheal an
c 26	25	17.0	6511	1 Q95493	Human Cdn-2 DNA. N
c 27	25	17.0	2072	1 Q95492	Human Cdn-1 cDNA.
c 28	25	17.0	1968	1 T17375	Bcl-1 cDNA. New is
c 29	25	17.0	2094	1 T42138	Bak gene. Screenin
c 30	25	17.0	2460	1 V44303	Human secreted pro
c 31	25	17.0	6623	1 T75251	Nucleotide sequenc
c 32	25	17.0	2094	1 V61498	Bak cDNA. New Bak-
c 33	24.8	16.9	1843	1 T38308	Human B-cell trans
c 34	24.6	16.7	2068	1 Q13224	DNA encoding Termi
c 35	24.6	16.7	1758	1 V58786	Human phospholipas
c 36	24.4	16.6	1621	1 Q56423	B. campestris Bp1
c 37	24.4	16.6	3300	1 V31822	Mutant Aspergillus
c 38	24.4	16.6	1855	1 V68998	DNA molecule encod
c 39	24.2	16.5	2035	1 N70687	DNA encoding human
c 40	24.2	16.5	2676	1 Q14850	Clone pT1284 enco
c 41	24.2	16.5	387	1 Q35904	Anti-CD4 V-lambda
c 42	24.2	16.5	1514	1 Q68267	Maize 2-acyltransf
c 43	24.2	16.5	2131	1 Q88155	Human lung tumour

Anti-HIV-1 Mab 447
Human G-protein ad

44 24.2 16.5 816 1 Q98724
45 24.2 16.5 2481 1 V33510

ALIGNMENTS

RESULT 1					
V32475					
ID	V32475 standard; cDNA; 1420 BP.				
AC	V32475;				
DT	11-SEP-1998 (first entry)				
DE	Bovine retinaldehyde binding protein cDNA.				
KW	Bovine retinaldehyde binding protein; retinal pigment epithelium; RPE;				
KW	11-cis-retinal; all-trans retinal; visual system; binding assay;				
OS	chromophore; ss.				
Bos sp.					
PH	Key	Location/Qualifiers			
FT	CDS	17..812			
FT		/*tag= a			
FT		/product= "Retinaldehyde binding protein"			
PN	US5763578-A.				
PD	09-JUN-1998.				
PF	16-DEC-1994; 358171.				
PR	16-DEC-1994; US-358171.				
PA	(FONG/) FONG H K W.				
PI	Fong HKW;				
DR	WPI; 98-347415/30.				
DR	P-PSDB; W48857.				
PT	Human and bovine retin-aldehyde-binding proteins - used to detect				
PT	aberration(s) of retinal binding in visual excitation systems				
PS	disclosure; Fig 1; 39pp; English.				
CC	The present sequence represents the bovine retinaldehyde binding				
CC	protein cDNA isolated from a bovine retinal pigment epithelium (RPE)				
CC	cDNA library. The bovine retinaldehyde binding protein binds both				
CC	11-cis-retinal and all-trans retinal. The invention claims that				
CC	molecular aberration of the visual system can be detected in binding				
CC	assays by observing any changes in the binding of the retinaldehyde				
CC	binding protein to its chromophores. The retinaldehyde binding protein				
CC	can also be used to raise antibodies, which in turn can be used to				
CC	detect changes of the protein in samples.				
Sequence	1420 BP; 291 A; 447 C; 384 G; 298 T;				
Query Match	18.48; Score 27; DB 1; Length 1420;				
Best Local Similarity	62.7%; Pred. No. 6.3;				
Matches 42; Conservative	0; Mismatches 25; Indels 0; Gaps 0;				
QY	7 ctggggccaacctgactcagtcctctctggaggccaacaggactctgagtcacctctgtgg 66				
Db	463 CTGGGGCCACTATGACTATGAGCCCTGGGGACCTGCTGCACCTGCACTATTCAGGGG 522				
QY	67 ggggtgga 73				
Db	523 GGACAGA 529				
RESULT 2					
Q25156					
ID	Q25156 standard; cDNA; 2158 BP.				
AC	Q25156;				
DT	18-NOV-1992 (first entry)				
DE	Alpha-GalNac from pAGB-3.				
KW	Lysosome; Schindler disease; infantile neuroaxonal dystrophy; ss.				
OS	Homo sapiens.				
PH	Key	Location/Qualifiers			
FT	CDS	345..1580			
FT		/*tag= a			
FT		/label= alpha-GalNac			
FT	signal_peptide	345..395			
FT		/*tag= b			
FT	mat_peptide	396..1580			
FT		/*tag= c			

FT poly_a_signal 2073. .2078
FT protein_bind /*tag= d
FT 2025. .2029
FT /*tag= e
FT /note= "recognised by the U4 small nuclear
FT ribonucleoprotein"
PN WO9207936-A.
PD 14-MAY-1991.
PF 23-OCT-1991; U07872.
PR 24-OCT-1990; US-602608.
PA (MOUN) MOUNT SINAI SCHOOL MEDICINE.
PI Bishop DF, Desnick RJ, Ioannou YA, Wang AM;
DR WPI; 92-183672/22.
DR P-PSDB; R24291.
PT Cloning and expression of alpha-n-acetyl-galactose aminidase -
PT used in enzyme replacement therapy for Schindler disease
PS Disclosure; Fig 2 (A-D); 71pp; English.
CC The sequence is of the PAB-3 cDNA insert contg. the complete coding
CC region for human alpha-GalNAC.
CC The availability of the full length cDNA for alpha-GalNAC allows
CC the study of the genomic organisation and evolution of this
CC lysosomal gene, and the characterisation of molecular lesions
CC causing Schindler disease.
SQ Sequence 2158 BP; 517 A; 610 C; 576 G; 455 T;

Query Match 18.2%; Score 26.8; DB 1; Length 2158;
Best Local Similarity 55.3%; Pred. No. 8.2;
Matches 52; Conservative 0; Mismatches 42; Indels 0; Gaps 0;

QY 32 ctgagggccacaggactctgagtcattctgtgggggtgggaggtgggacaggaagg 91
DB 1542 CTGTATCCCATCAAGAACCTGGAGATGTCACAGCAGTGGGACATGTCACAGG 1601
QY 92 ggtgaatggtactgctgattacaacctctgtgtgc 125
DB 1602 CTGTGTGGCACCCTGAGCCTAGACCATGGAGC 1635

RESULT 3
ID Q81826 standard; cDNA; 1840 BP.
AC Q81826;
DE 10-MAR-1995 (first entry)
DE Alpha-N-acetylgalactosaminidase.
KW erythrocyte; amplification; primer; polymerase chain reaction; PCR;
KW probe; blood; type A; type B; type AB; type O; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT cds 73..1308
FT /*tag= a
PN WO9411518-A.
PD 26-MAY-1994.
PF 08-NOV-1993; U10794.
PR 18-NOV-1992; US-977945.
PA (GENW) GENENCOR INT INC.
PI Berka RM;
DR WPI; 94-183517/22.
DR P-PSDB; R69101.
PT Prodn. of human placental alpha-N-acetylgalactosaminidase - by
PT expression in transformed host cells, used to convert type A, B
PT or AB erythrocytes to type O
PS Claim 13; Fig 2; 66pp; English.
CC The two primers given in Q81823-24 were used in the screening of
CC libraries contg. sequences specific for alpha-N-
CC acetylgalactosaminidase cDNA clones. A 466 bp fragment was
CC obtained, which was then subcloned and used as a probe.
CC Another probe given in Q81825 was used in a secondary
CC screening process. A full length alpha-N-acetylgalactosaminidase
CC cDNA was obtained (see Q81826).
SQ Sequence 1840 BP; 417 A; 539 C; 485 G; 399 T;

Query Match 18.2%; Score 26.8; DB 1; Length 1840;
Best Local Similarity 55.3%; Pred. No. 7.9;
Matches 52; Conservative 0; Mismatches 42; Indels 0; Gaps 0;

QY 32 ctgagggccacaggactctgagtcattctgtgggggtgggaggtgggacaggaagg 91
DB 1270 CTGTATCCCATCAAGAACCTGGAGATGTCACAGCAGTGGGACATGTCACAGG 1329
QY 92 ggtgaatggtactgctgattacaacctctgtgtgc 125
DB 1330 CTGTGTGGCACCCTGAGCCTAGACCATGGAGC 1363

RESULT 4
ID T28078/c standard; cDNA to mRNA; 334 BP.
AC T28078;
DE 16-OCT-1996 (first entry)
DE Human gene signature HUMGS08314.
KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;
KW human; cloning; mapping; non-biased library; diagnosis; detection;
KW cell typing; abnormal cell function; ss.
OS Homo sapiens.
FH WPI; 95-206931/27.
PN WO9514772-A1.
PD 01-JUN-1995.
PF 11-NOV-1994; J01916.
PR 12-NOV-1993; JP-355504.
PA (MATS/) MATSUBARA K.
PA (OKUB/) OKUBO K.
PI Matsubara K, Okubo K;
DR WPI; 95-206931/27.
DE Identifying gene signatures in 3'-directed human cDNA library - e.g.
DE for diagnosis of abnormal cell function, by preparing cDNA that
DE reflects relative abundance of corresp. mRNA in specific human
DE tissues
PS Claim 1; Page 1996; 2245pp; Japanese.
CC A single-stranded DNA (or its complementary strand or the corresp.
CC double-stranded DNA) which comprises one of the 7837 "GS" sequences
CC given in T19001-T26837 and which is able to hybridise to part of
CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
CC sequences were obtained from 3'-directed cDNA libraries prepared
CC from various human tissues; synthesis of cDNA was initiated from the
CC 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
CC untranslated sequence is unique to a particular mRNA species, almost
CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
CC is constructed so as to reflect accurately the relative abundance of
CC different mRNAs in the particular tissue from which it was derived.
CC The appearance frequency of a given GS in a cDNA library can be
CC determined (esp. using primers and probes derived from the GS
CC sequences) as a means of diagnosing abnormal cell function or for
CC recognising different cell types.
SQ Sequence 334 BP; 65 A; 98 C; 79 G; 79 T;

Query Match 18.2%; Score 26.8; DB 1; Length 334;
Best Local Similarity 63.5%; Pred. No. 5;
Matches 40; Conservative 0; Mismatches 23; Indels 0; Gaps 0;

QY 37 ggcacacaggactctgagtcattctgtgggggtgggaggtgggacaggaagggtga 96
DB 127 GGACACTCTGACTATTGGACAAATCTCTGGGGAGTNCCTCCAGGGGAAGGAGGTGT 68
QY 97 atg 99
DB 67 GTG 65

RESULT 5
ID N91109 standard; DNA; 495 BP.
AC N91109;
DT 21-JUN-1990 (first entry)


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CC sequences of the probes can also be used for the detection of the
CC bacteria [sic] associated with periodontal disease. The probes can detect
CC the presence of periodontopathic bacteria. The probes and primers provide
CC a quick and specific identification without a need for the initial
CC knowledge of the sequence of the targetted DNA. 44 T;
SQ Sequence 142 BP; 31 A; 34 C; 31 G; 44 T;

Query Match 17.7%; Score 26; DB 1; Length 142;
Best Local Similarity 70.0%; Pred. No. 7.2;
Matches 35; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 60 cctgtgggggtggaggtgggacaaagggaagggtgaatggtactgtcta 109
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 113 CCTGCGGGTGGAAAGGCAAGGTAGGGGGTGGCAGGCACGACAGA 64

RESULT 11
V02016/c
ID V02016 standard; cDNA to mRNA; 2525 BP.
AC V02016;
DE 18-JUN-1998 (first entry)
KW DNA encoding tumour antigen protein.
KW Tumour antigen protein; gastric cancer; intracellular digestion;
KW bind; major histocompatibility complex class I antigen; recognition;
KW T-cell; gene therapy; tumour; autoimmune disease; ds.
OS Homo sapiens.
FH Key Location/Qualifiers
FT 5'UTR 1..38
FT CDS /*tag= a
FT 3'UTR 39..2441
FT polyA_signal /*tag= b
FT /*tag= c
FT /*tag= d
FT /*note= "polya site"
PN W09746676-Al.
PD 11-DEC-1997.
PR 04-JUN-1997; J01893.
PR 25-NOV-1996; JP-330424.
PR 07-JUN-1996; JP-168429.
PR 08-OCT-1996; JP-287572.
PR (Itoh/) ITOH K.
PI Imai Y, Itoh K, Shichijo S;
DR WPI; 98-042184/04.
DR P-PSDB; W44003.
PT DNA encoding tumour antigen protein, fragments of which bind to MHC
PT class I antigens - useful in gene therapy and auto-immune diseases
PS Claim 2; Pages 40-42; 49pp; Japanese.
CC The present sequence encodes a tumour antigen protein. It was isolated
CC from a gastric cancer cell line. The tumour antigen protein has the
CC ability to form fragments by intracellular digestion which bind to major
CC histocompatibility complex (MHC) class I antigens to form a complex
CC which is recognised by T-cells. The DNA is useful for gene therapy of
CC tumours and autoimmune diseases.
CC Sequence 2525 BP; 625 A; 653 C; 906 G; 341 T;

Query Match 17.6%; Score 25.8; DB 1; Length 2525;
Best Local Similarity 53.5%; Pred. No. 18;
Matches 54; Conservative 0; Mismatches 47; Indels 0; Gaps 0;

QY 45 ggaactctgagcatcctgtgggtggaggtgggacaaagggaagggtgaatggtact 104
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 2381 GGTCTCTGAGCTTCTGCTTCTGCTGAGCAGGCGCCAGGGGGGTGTCGT 2322

QY 105 gctgtatacaacctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtttat 145
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 2321 GGAGCTCACTTCTTCTCAGGAGCGCTCTCTCGTCCAGCTTCT 2281

RESULT 12

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Q61216
ID Q61216 standard; DNA; 383 BP.
AC Q61216;
DE 16-MAR-1994 (first entry)
DE Human brain Expressed Sequence Tag EST01251.
KW Gene transcription product; genetic markers; tagging; in vivo;
KW transcription; mapping; locations; chromosomes; chromosomal; ss.
OS Homo sapiens.
PN W09316178-A.
PD 19-AUG-1993.
PR 12-FEB-1993; U01294.
PR 12-FEB-1992; US-837195.
PA (USSH ) US DEPT HEALTH & HUMAN SERVICE.
PI Adams MD, Moreno RF, Venter CJ;
DR WPI; 93-272882/34.
PT Enriched oligonucleotides and corresp. sequences - used as
PT markers for human genes transcribed in-vivo, facilitate tagging
PT of most human genes
PS Example 4; Page 456; 500pp; English.
CC The Expressed Sequence Tag was isolated from a human brain cDNA
CC library as part of a large set of ESTs which can be used as markers
CC for human genes transcribed in vivo. They can be used to facilitate
CC tagging of most human genes, for mapping locations of expressed genes
CC on chromosomes, for individual or forensic identification, for mapping
CC locations of disease-associated genes, for identification of tissue
CC type, and for prepn. of antisense sequences, probes and constructs.
CC EST01251 has a "good" coding probability as evaluated using the
CC coding-region prediction program CRM. See also Q59041-Q61440.
SQ Sequence 383 BP; 118 A; 90 C; 94 G; 80 T;

Query Match 17.4%; Score 25.6; DB 1; Length 383;
Best Local Similarity 54.2%; Pred. No. 13;
Matches 52; Conservative 0; Mismatches 44; Indels 0; Gaps 0;

QY 12 ccaaccatgactcagtgctctgttgaggaggaagggtgaatggtactgct 107
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 169 CCTACTTTGGCGGTCTATACTGAAGCCCTCCACTCCCTCCTCAAGAGGTGAG 228

QY 72 gaggtgggacaaagggaagggtgaatggtactgct 107
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 229 GAGGTGGGAGAGAGCAGTGAGGAGCAAGATATGCT 264

RESULT 13
Q58718
ID Q58718 standard; DNA; 1254 BP.
AC Q58718;
DE 30-SEP-1994 (first entry)
DE Human gamma-1 chain first membrane exon.
KW Membrane; exon; human; gamma-1; chain; membrane anchoring peptides;
KW heavy chain; isotype; cell surface; extracellular region;
KW mlgis segment; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT Key 547..678
FT exon /*tag= a
FT US298420-A.
PN 29-MAR-1994.
PD 03-AUG-1990; 562201.
PR 03-AUG-1990; US-562201.
PR 19-JUN-1992; US-902449.
PA (TANO-) TANOX BIOSYSTEMS INC.
PI Chang TW;
DR WPI; 94-100338/12.
DR P-PSDB; R60128.
PT Monoclonal antibodies and fragments specific for extracellular
PT mlgis epitope(s) on B cells - used to bind B cells for
PT immunosuppressive purposes or to remove B cells from the
PT circulation
PS Disclosure; Fig 1A; 13pp; English.
CC The sequences given in Q58718-19 represent the first and second
CC membrane exons of the human gamma-1 chain which encode membrane

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